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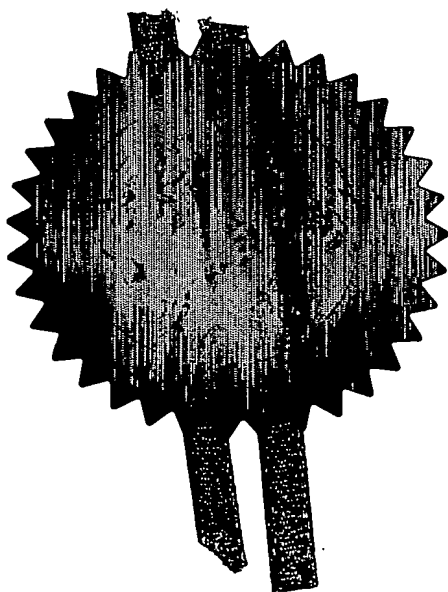
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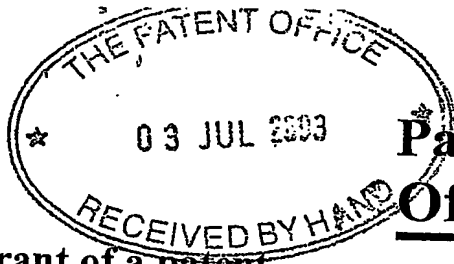
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1. Your reference

P511586GB /50440

2. Patent application number

(The Patent Office will fill in this part)

0315629.6

3 JUL 2003

3. Full name, address and postcode of the or of each applicant (underline all surnames)

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Patents ADP number (if you know it)

8604053001

If the applicant is a corporate body, give the country/state of its incorporation

United Kingdom

4. Title of the invention

New Uses for Antibiotic

5. Name of your agent (if you have one)

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

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Patents ADP number (if you know it)

1776001

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
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Date of filing
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7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

a) any applicant named in part 3 is not an inventor, or
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YES

See note (d))

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Description 13

Claim(s) 2

Abstract

Drawing (s)

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77)

Request for substantive examination (Patents Form 10/77)

Any other documents (please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature *Wethers & Rogers* Date *3rd July 2003*

12. Name and daytime telephone number of person to contact in the United Kingdom

Adrian Tombling

020 7663 3500

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New Uses for an Antibiotic

The present invention relates to the use of antibiotics as medicaments for the treatment or prophylaxis of disease and infections especially in pigs and poultry.

Pigs and poultry, especially those which are intensively reared or reared in large-scale operations, have a tendency to suffer from or risk catching a variety of diseases and infections, for example, *Brachyspira pilosicoli* in pigs and *Ornithobacterium rhinotracheale* in poultry.

Brachyspira pilosicoli causes a condition known as spirochetel diarrhoea, which features diarrhoea and slow growth. The infection rarely causes death, but can be fatal, particularly when associated with infection with another intestinal parasite. Infection can have a major economical impact on the profitability of pig production due to reductions in daily weight gain and feed conversion. Unlike some other porcine intestinal infections *Brachyspira pilosicoli* is not solely associated with pigs. It is also known to infect humans, dogs and other animals.

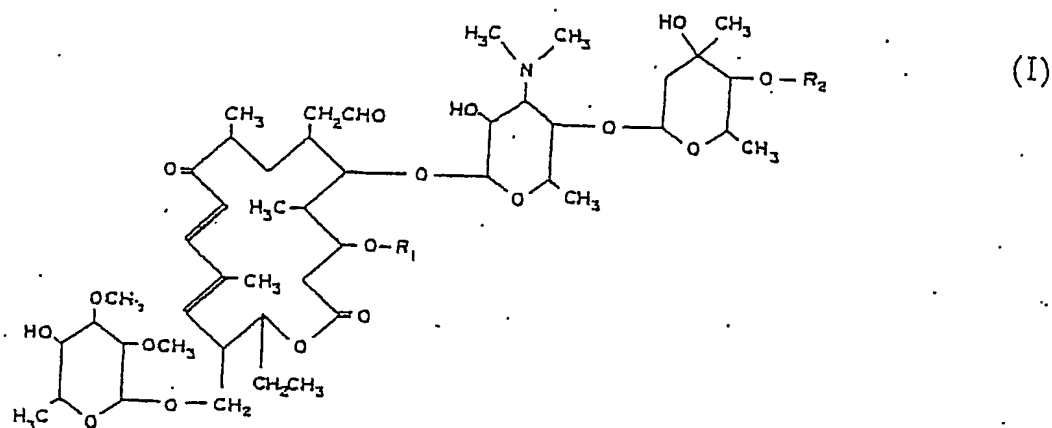
Ornithobacterium rhinotracheale is a respiratory disease characterised by mild respiratory symptoms, increased mortality and poor weight gain and feed conversion. It causes economic losses due to increased medical costs, reduced growth and high condemnation rates at processing. *Ornithobacterium rhinotracheale* has been isolated from many species including chicken, duck, partridge, goose, pigeon, and turkey, amongst others. This indicates there is a broad potential reservoir.

Although there are some known treatments for *Brachyspira pilosicoli* and *Ornithobacterium rhinotracheale*, they are mostly ineffective due to widespread resistance to commonly used antibiotics.

Surprisingly, the inventors have found that the known antibiotic aivlosin (otherwise known as 3-O-acetyl-4"-O-isovaleryl-tylosin), which has previously been used in high

doses for the treatment and control of *Mycoplasma* diseases in poultry, is also effective in the prevention and treatment of *Brachyspira pilosicoli*, particularly in pigs and of *Ornithobacterium rhinotracheale* particularly in poultry.

In British Patent Specification No. 1,539,907 there are disclosed tylosin derivatives having acyl groups in the 3 and 4" positions and acid addition salts thereof, specifically the tartaric, acetic, propionic, citric, succinic, hydrochloric, sulphuric and phosphoric acid addition salts. Amongst the tylosin derivatives specifically disclosed there is 3-O acetyl-4"-O-isovaleryl-tylosin, which is now commonly known as aivlosin. This compound has the formula



where R_1 is acetyl and R_2 is isovaleryl. There is also disclosed a process for the production of aivlosin by the biochemical acylation of tylosin or an appropriately partially acylated tylosin by means of an appropriate acylating micro-organism of the genus *Streptomyces*, especially one selected from *Streptomyces thermotolerans* (ATCC 11416), *Streptomyces fungicidus* subsp. *espinomyceticus* (ATCC 21574), *Streptomyces mycarofaciens* (ATCC 21454) and *Streptomyces hygroscopicus* (ATCC 21582), in the presence of the appropriate acyl donor, especially acetyl CoA, isovaleryl CoA, acetic

acid, isovaleric acid, potassium, sodium or ammonium salts of those acids, methanol and ethanol esters of these acids, amides of these acids and α -oxovaleric acid.

British Patent Specification No. 1,539,907 mentions that the tylosin derivatives can be administered to humans or animals and refers to their activity against a number of gram-positive bacteria, including some drug-resistant bacteria, but it does not specifically refer to the use of the derivatives in the treatment or control of specific diseases or infections of animals, although it does say that they can be employed on humans, livestock, household pets, laboratory animals and poultry and in the enteral, parenteral or topical control of infectious diseases in a similar manner as for known macrolide antibiotic drugs.

In fact, aivlosin on the basis of its initial Japanese marketing registration (No 4 chika AC1771) has to date been marketed and approved for marketing only for the treatment and control of Mycoplasma diseases in pigs and poultry at high doses of 200 to 500 ppm in feed. There should be no reason to suppose that it would be suitable for the treatment and prophylaxis of other infections and diseases, particularly of pigs and poultry. Other macrolide antibiotics having an effectiveness against mycoplasma diseases such as erythromycin do not have any effect or any significant effect against other infections of pigs and poultry such as those mentioned above. It is, of course, a feature of the approvals schemes which apply in all major countries that a veterinary medicament which is approved for marketing for one specific purpose cannot be marketed or recommended for use for any other specific purpose without a separate authorisation or approval from the relevant authority. There is thus a strong counter-incentive to the research into the use of even known antibiotics for new veterinary uses.

PCT application WO 02/32233 describes the use of aivlosin to treat *Brachyspira hyodysenteriae* in pigs. *Brachyspira hyodysenteriae* is an infection of the large intestine. It causes swine dysentery, resulting in bloody diarrhoea and death through dehydration. The activity of aivlosin against *Brachyspira pilosicoli* could not have been predicted

from the use against *Brachyspira hyodysenteriae* because the efficacy of any antimicrobials cannot be accurately defined without tests on live animals.

WO 02/32233 also discloses the use of aivlosin to treat *Lawsonia intracellularis* in pigs. *Lawsonia intracellularis* is a disease of the small intestine which causes diarrhoea and wasting. *Lawsonia intracellularis* can be categorised as a Gram negative organism, but it is not a traditional Gram negative bacterium.

Aivlosin is characterised as a macrolide antibiotic, effective against Gram positive bacteria and mycoplasma. Such antibiotics are not expected to be effective against Gram negative bacteria (Antimicrobial Therapy in Veterinary Medicine, 3rd Ed. (2000) Edited by Prescott JF, Baggot JD and Walker RD. Iowa State University Press), as demonstrated by existing data of the use of macrolides against *Pasteurella haemolytica*. Activity of macrolides cannot be predicted from the bacterial wall structure. Macrolides have been demonstrated to be effective against both *Mycobacterium* spp., the cell walls of which incorporate complex lipid conjugates and the wall-less *Mycoplasma* spp. The activity of any macrolide cannot be predicted without specific testing. Against expectation, the inventors have found that it is also effective against the Gram negative bacteria *Ornithobacterium rhinotracheale*.

From extensive *in vitro* and *in vivo* (animal) trial work, the inventors have confirmed that aivlosin and acceptable derivatives thereof are effective in the prevention and treatment of *Brachyspira pilosicoli* in pigs and *Ornithobacterium rhinotracheale* in poultry at reasonable dose rates.

The present invention provides for the use of aivlosin, as such or as a pharmacologically acceptable (non-toxic) derivative, such as an acid addition salt, in the preparation of a medicament for the treatment or prophylaxis of *Brachyspira pilosicoli* and *Ornithobacterium rhinotracheale* in animals. The medicament is preferably for treating *Brachyspira pilosicoli* in mammals, more preferably in pigs. In an alternative embodiment the medicament is also for treating *Ornithobacterium rhinotracheale* in birds, more preferably in poultry. Also provided is a method of

treatment or control of *Brachyspira pilosicoli* and *Ornithobacterium rhinotracheale* in animals comprising administering to an animal an effective amount of aivlosin or a pharmacologically effective derivative thereof.

The term "pigs" encompasses all members of the pig family, for example, members of the *Suidae* family. The term "poultry" encompasses all types of domestic fowl, including, but not limited to chickens, turkey, ducks and geese.

Preferably the medicament for the treatment or prophylaxis of *Brachyspira pilosicoli* in pigs is added to food at a level of 10 to 200 ppm, more preferably 10 to 100 ppm, even more preferably 20 to 50 ppm.

Preferably the medicament for the treatment or prophylaxis of *Ornithobacterium rhinotracheale* in poultry is added to food at a level of 10 to 200 ppm, more preferably 10 to 100 ppm, even more preferably 20 to 50 ppm.

Either medicament is preferably suitable for addition to food or drinking water. Alternatively the medicament may be suitable for administration by injection.

Also provided by the invention is the use of aivlosin to prevent or reduce growth of *Brachyspira pilosicoli* or *Ornithobacterium rhinotracheale* *in vitro*. The prevention or reduction of growth of these bacteria can be useful in the *in vitro* preparation of intestinal tissues, or in the comparison of the activity of antibiotics against bacteria.

Aivlosin is available in free form as a white crystalline powder having a melting point of 180°C-184°C, soluble in lower alcohols such as ethanol, ketones such as acetone, ethers such as diethyl ether, esters such as ethyl acetate and aromatic hydrocarbons such as toluene, although it is barely soluble in n-hexane and petroleum ether. It is very soluble in aqueous solutions of pH around and below 7 but less soluble in aqueous solutions of higher pH. Because it is a basic compound it forms acid addition salts, and the use of such salts which are pharmacologically acceptable is also included within the present invention. Acids to form acceptable acid addition salts include inorganic acids

such as hydrochloric, sulphuric or phosphoric acid and organic acids such as tartaric, acetic, propionic, citric and succinic acids. Specific examples of acceptable derivatives are aivlosin hydrochloride (melting point 129-133°C) and aivlosin tartrate (melting point 119-122°C). Such derivatives are frequently more water-soluble than aivlosin itself and their use may therefore have formulation advantages.

Derivatives of aivlosin preferably include any pharmacologically acceptable functional derivatives. The functional derivatives may be produced by modifying one or more of the substituent groups of aivlosin. Preferably the derivative is a salt; most preferably an acid salt.

Aivlosin and appropriate derivatives can be formulated according to the present invention into medicaments in known ways, for example to provide compositions for oral, enteral or parenteral administration, by admixing with appropriate solid or liquid carriers and excipients for the administration route desired. Conventional ingredients can be used as carriers and excipients, for example water and salt solutions for liquid formulations and siliceous materials-silica and silicates (such as hydrated magnesium silicate), cereal products (such as soybean meal and wheat flour) and other pharmacologically acceptable solids for solid formulations for oral administration. The formulations can also contain further auxiliaries and additives such as minerals, lubricants, preservatives, stabilisers, wetting agents, emulsifiers, buffers and colouring or flavouring materials in a conventional manner. In the prophylaxis or control of the diseases mentioned it is particularly convenient to include the aivlosin or derivative as an additive to animal feed or drinking water for animals, but in the treatment of the disease it can be included in an injectable solution, or a tablet, capsule or syrup, if desired.

Aivlosin (as such or in the form of an appropriate derivative, for example an acid addition salt such as the tartrate) may be formulated into premixes in various potencies from 1 to 10% by weight. A particularly suitable composition for producing such premixes comprises aivlosin salt, filler such as soybean powder and additives such as hydroxypropyl cellulose and has a potency of 180 to 220 mg/g.

In order to ensure stability of aivlosin in animal feed which may have been subjected to high-temperature processing for pelleted or extruded feed it is desirable to provide a coated aivlosin (as such or in the form of an appropriate derivative, for example an acid addition salt such as the tartrate) in particulate form coated with polyvinylpyrrolidone. Suitable proportions by weight are in the range of 50:1 to 1:1 active ingredient: polyvinylpyrrolidone. Inert fillers and other ingredients may be present in such compositions, the overall polyvinylpyrrolidone concentration being preferably 0.1 to 10% by weight.

The medicament formulations for use either as feed additives or as directly administered preparations may contain any convenient proportion of aivlosin for example from 1% or less to 90% or more, by weight. Liquid formulations typically contain 50 to 90% by weight, whereas solid formulations typically contain 1 to 25% by weight.

For treatment or control of *Brachyspira pilosicoli* infections in pigs aivlosin may for example be administered in feed at a rate of 10 to 200 ppm by weight (10 to 200 g per 1,000 kg of feed) for a period of time long enough to control or treat the disease successfully, for example 7-14 days.

For treatment or control of *Ornithobacterium rhinotracheale* infections in poultry aivlosin may for example be administered in feed at a rate of 20 to 50 ppm by weight (20 to 50g per 1,000 kg of feed) for a period of time long enough to control or treat the disease successfully, for example 7-14 days. Alternatively aivlosin may be administered at a rate of between 100 to 150 ppm by weight in drinking water (100 to 150g) per 1000 l of water).

The following examples, in which parts are by weight, illustrate the use of aivlosin in the manufacture of veterinary medicaments or preparations for treatment or prophylaxis of the animal infections according to the present invention.

Example 1

20 parts of aivlosin API (active pharmaceutical ingredient) made into a solution in water is mixed with 80 parts of soybean meal, and the mixture is spray dried to give a solid additive for feedstuff containing 200 kg aivlosin activity per 1000 kg. This formulation can be added to pig and poultry feed to provide an in-feed concentration of aivlosin of 25 to 200 g aivlosin per 1000 kg final feed.

Example 2

25 parts of aivlosin 20% is mixed with 50 parts of hydrated magnesium silicate (an inert silica), 24 parts of wheat feed flour and 1 part of liquid paraffin EP as a powder blend to give a solid additive for feedstuff containing 50 kg aivlosin activity per 1000 kg. This formulation can be used in pig and poultry feed as in Example 1.

Example 3

5 parts of aivlosin 20% as used in Example 2 is mixed with 40 parts of hydrated magnesium silicate, 54 parts of wheat feed flour and 1 part of liquid paraffin EP as a powder blend to give a solid additive for feedstuff containing 10 kg aivlosin activity per 1000 kg. This formulation can be used in pig and poultry feed as in Example 1.

Example 4

Aivlosin is dissolved in water to provide an aqueous solution containing 80-90% aivlosin activity for use in drinking water for pigs or poultry. This formulation can be added to drinking water to provide aivlosin concentrations in drinking water in the range 25 to 100 g per 200 litres of drinking water.

Example 5

Aivlosin API containing more than 80% w/w aivlosin tartrate was mixed into an 850 kg batch comprising

Aivlosin API	163-169 kg
Hydroxypropyl cellulose. Ph. Eur.	8.2-8.5 kg
Water, Ph. Eur	800-1200 litres
Non-fat soybean powder	720 kg

The batch was processed and the water was removed during processing. The input of aivlosin API was adjusted for content value of free base, determined by HPLC, of the raw material to achieve a final product bioassay potency of 180-220 mg/g. The product (AIVLOSIN FG 200), which could also be produced in other batch sizes, was suitable for manufacturing aivlosin premixes in various potencies from 1% to 10%.

Example 6

Coated aivlosin formulations possessing stability in animal feed after high-temperature processing for pelleted or extruded feed were produced in batches of 1000 kg (although other batch sizes could be used) from the following ingredients:

AIVLOSIN FG 200 (see Example 5)	250.0 kg
Paraffin, Light Liquid, Ph. Eur.	10.0 kg
Wheat feed flour	240.0 kg
Polyvinylpyrrolidone	10.0 kg - 100.0 kg
Sepiolite	to 1000.0 kg

Example 7

Determination of Minimum Inhibitory Concentration (MIC) of Aivlosin against *Brachyspira pilosicoli*.

The MIC of acetyl isovaleryltylosin was determined against 5 field strains of *Brachyspira pilosicoli* isolated from different pig populations in various parts of England using an antibiotic dilution method. Four replicates of 0.2 ml of each isolate were inoculated into agar plates prepared with antibiotic concentrations from 0.78 to 200 µg/ml. A strain of *Brachyspira hyodysenteriae* (P18A), for which the MIC for acetyl isovaleryltylosin was already known, was used as a control. The determined MICs are given in the table below.

Strain Ref.	MIC (µg/ml)
P0098-10-97	25.0
P0352-10-98	6.25
P0730-09-97	12.5
P0204-10-97(4)	<0.781
P0525-01-98	6.25
Control P18A	12.5

It is concluded that the MICs for acetyl isovaleryltylosin were generally similar, falling in the range 6.25-25.0 µg/ml with the exception of one isolate (P0204-10-97 (4)) against which acetyl isovaleryltylosin was far more active.

The results show that aivlosin is particularly effective at preventing the growth of *Brachyspira pilosicoli*, even at relatively low concentrations.

Example 8

Testing of aivlosin and tilmicosin (pulmotil) against *Ornithobacterium rhinotracheale* using the MIC (minimum inhibitory concentration) test.

The following antibiotics were each tested against four isolates of *Ornithobacterium rhinotracheale* (OR):

- 1) Aivlosin
- 2) Pulmotil

The antibiotics were tested against the following isolates of OR:

- 1) 568/99
- 2) 587/00
- 3) 33/01
- 4) 1322/01

- 1) MIC method

The concentration needed of each antibiotic was calculated according to its active ingredient.

Aivlosin

81% active

$$32 \mu\text{g/ml} \times 2 = 64 \mu\text{g/ml}$$

$$64 \mu\text{g/ml} \times 1.23 = 78.72 \mu\text{g/ml}$$

$$= 0.04 \text{ g/500 ml}$$

Pulmotil

25% active

$$32 \mu\text{l/ml} \times 2 = 64 \mu\text{l/ml}$$

$$64 \mu\text{l/ml} \times 4 = 256 \mu\text{l/ml}$$

$$= 0.13 \text{ ml/500 ml}$$

The four freeze dried bacterial isolates of *Ornithobacterium rhinotracheale* were reconstituted, inoculated into 10 ml serum broth (supplied by OBP, code 655) and incubated at 37°C for 48 hours. After incubation the optical density (OD) of each culture was read at 540 nm. To verify purity the cultures were plated out onto Blood

Tryptose agar plates, incubated at 37°C, using 5 % CO₂ conditions, for 48 hours after which the plates were examined for any contaminants.

12 tubes per isolate for each of the antibiotics were placed into a test tube rack. 2 ml serum broth was dispensed into each of the tubes. 2 ml of the antibiotic were then added to the first tube in the respective row and a twofold dilution were made (32µg/ml – 0.0625µg/ml). 20 µl of the bacteria was then added to 11 of the tubes. The 12th tube, i.e. negative control received neither antibiotic nor bacteria. The 11th tube i.e. positive control only received 20 µl bacteria. All the tubes were then incubated at 37°C and the MIC read after 48 hours incubation

2) Results

Optical density readings (540 nm) before addition to antibiotic dilutions :

- 1) 568/99 – 1.025
- 2) 587/00 – 1.045
- 3) 33/01 – 1.058
- 4) 1322/01 – 1.081

1) Aivlosin

Dilution (µg/ml)	Bacterial Growth			
	568/99	587/00	33/01	1322/01
32	-	-	-	-
16	-	-	-	-
8	-	-	-	-
4	-	-	-	-
2	-	-	-	-
1	-	-	-	-
0.5	-	-	-	-
0.25	+	+	+	+
0.125	+	+	+	+
0.0625	+	+	+	+
Negative control	-	-	-	-
Positive control	+	+	+	+

2) Pulmotil

Dilution ($\mu\text{g/ml}$)	Bacterial Growth			
	568/99	587/00	33/01	1322/01
32	-	-	-	-
16	-	-	-	-
8	-	-	-	-
4	-	-	-	+
2	+	+	+	+
1	+	+	+	+
0.5	+	+	+	+
0.25	+	+	+	+
0.125	+	+	+	+
0.0625	+	+	+	+
Negative control	-	-	-	-
Positive control	+	+	+	+

3) Conclusions :

A concentration of 0.5 $\mu\text{g/ml}$ Aivlosin inhibited the growth of the isolates of *Ornithobacterium rhinotracheale* tested.

A concentration of 4 $\mu\text{g/ml}$ Pulmotil inhibited the growth of three of the isolates of *Ornithobacterium rhinotracheale* tested. Inhibition of the fourth isolate was only achieved at a concentration of 8 $\mu\text{g/ml}$.

The results show that aivlosin is particularly effective at preventing the growth of *Ornithobacterium rhinotracheale*, even at relatively low concentrations.

Claims

1. The use of aivlosin or a pharmacologically acceptable derivative thereof for the preparation of a medicament for the treatment or prevention of a *Brachyspira pilosicoli* infection.
2. The use of claim 1 wherein the medicament is for treating pigs.
3. The use of claim 1 or 2 wherein the medicament is for addition to feed at a rate of 10 to 200 ppm.
4. The use of any of claims 1 to 3 wherein the medicament is an additive for feed or drinking water.
5. A method of treating or preventing a *Brachyspira pilosicoli* infection comprising administering to an animal an effective amount of aivlosin or a pharmacologically acceptable salt thereof.
6. The method of claim 5 wherein the animal is a pig.
7. The method of claim 5 or claim 6 wherein the aivlosin or a pharmacologically acceptable derivative thereof is administered in either feed or drinking water.
8. The use of aivlosin or a pharmacologically acceptable derivative thereof for the preparation of a medicament for the treatment or prevention of an *Ornithobacterium rhinotracheale* infection.
9. The use of claim 8 wherein the medicament is for treating poultry.
10. The use of claims 8 or 9 wherein the medicament is for addition to feed at a rate of 10 to 200 ppm.

11. The use of any of claims 8 to 10 wherein the medicament is an additive for feed or drinking water.

12. A method of treating or preventing an *Ornithobacterium rhinotracheale* infection in animals comprising administering to an animal an effective amount of aivlosin or a pharmacologically acceptable derivative thereof.

13. The method of claim 12 wherein the animal is a poultry bird.

14. The method of claim 13 wherein the aivlosin or a pharmacologically acceptable derivative thereof is administered in either feed or drinking water.

15. The use of aivlosin or a pharmacologically acceptable derivative thereof in the prevention or reduction of growth of *Brachyspira pilosicoli* *in vitro*.

16. The use of aivlosin or a pharmacologically acceptable derivative thereof in the prevention or reduction of growth of *Ornithobacterium rhinotracheale* *in vitro*.

PCT/GB2004/002887

